

**BYOMA ACNE TREATMENT SPOT- salicylic acid paste**  
**Byoma Ltd**

-----  
**Byoma Acne Treatment Spot Paste**

**Drug Facts**

**Active Ingredients**

Salicylic Acid 2%

**Purpose**

Acne Treatment

**Use**

- for the treatment of acne

**Warnings**

**For external use only**

**When using this product**

- skin irritation and dryness is more likely to occur if you use another topical acne medication at the same time. If irritation occurs, only use one topical acne medication at a time.

**Stop use and ask a doctor if**

- skin irritation occurs or gets worse.

**Keep out of reach of children.**

If swallowed, get medical help or contact a Poison Control Centre right away.

**Directions**

- Clean the skin thoroughly before applying this product
- Cover the entire affected area with a thin layer one to three times daily.
- Because excessive drying of the skin may occur, start with one application daily, then gradually increase to two or three times daily if needed or as directed by a doctor.
- If bothersome dryness or peeling occurs, reduce application to once a day or every other day.

**Other Information**

Protect the product in this container from excessive heat and direct sun.

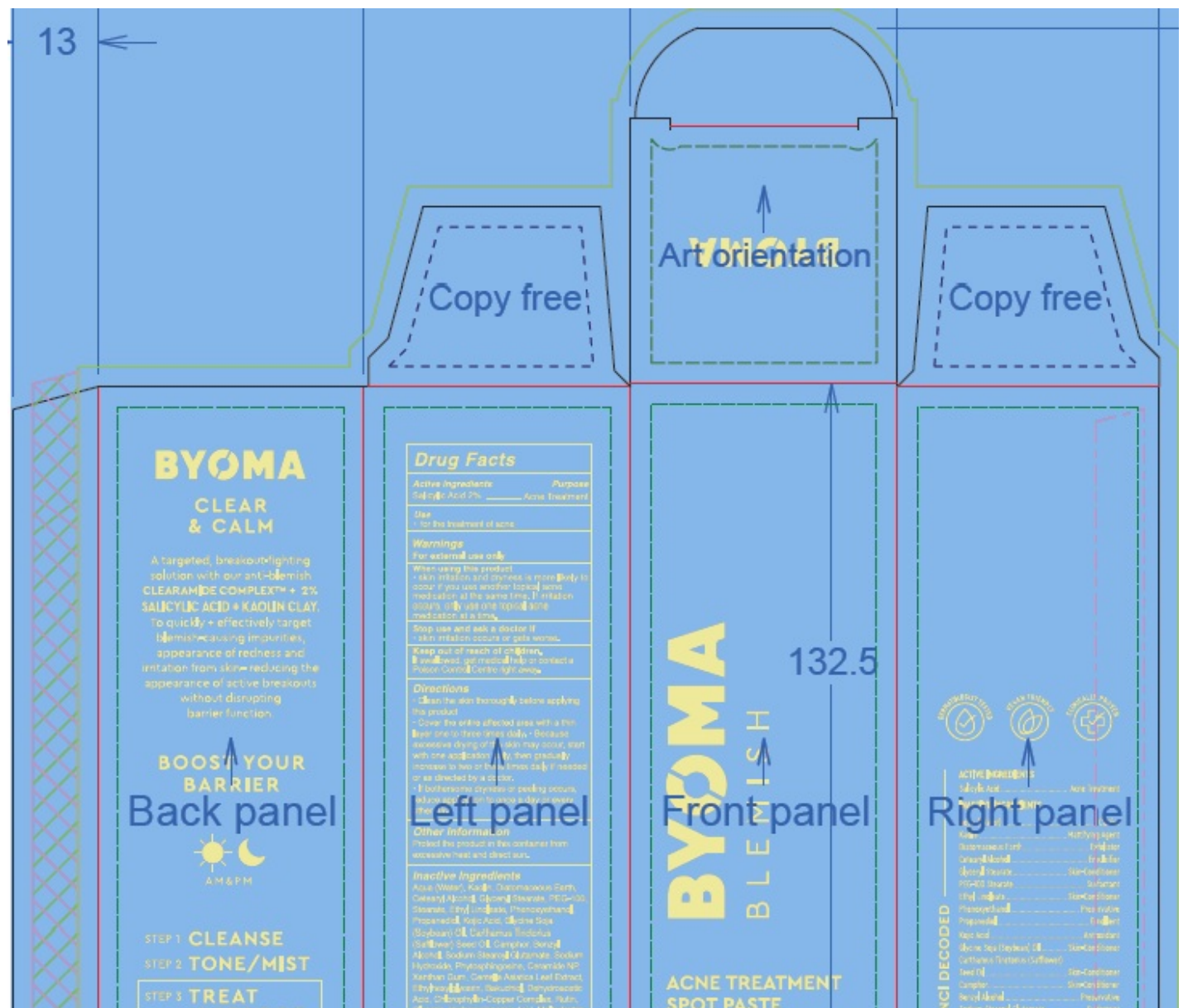
## Inactive Ingredients

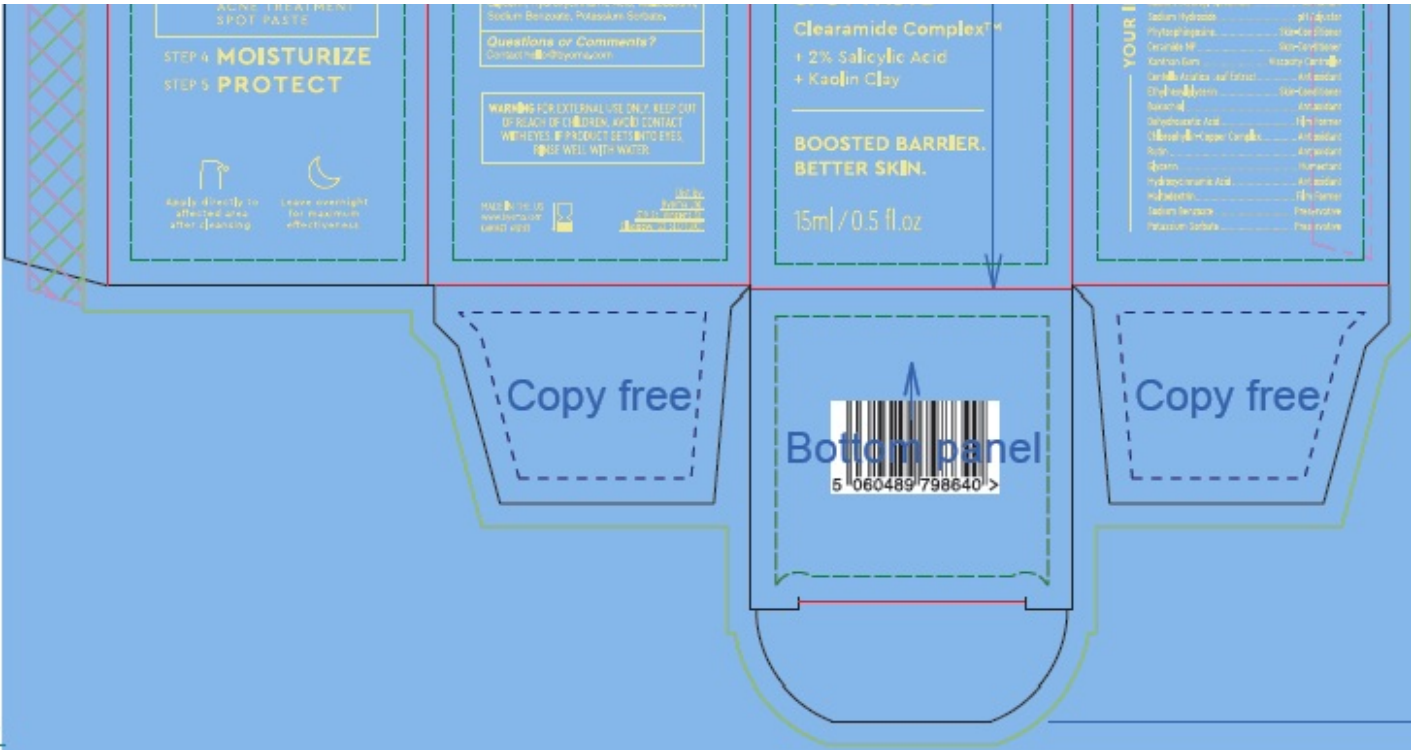
Aqua (Water), Kaolin, Diatomaceous Earth, Cetearyl Alcohol, Glyceryl Stearate, PEG-100 Stearate, Ethyl Linoleate, Phenoxyethanol, Propanediol, Kojic Acid, Glycine Soja (Soybean) Oil, Carthamus Tinctorius (Safflower) Seed Oil, Camphor, Benzyl Alcohol, Sodium Stearoyl Glutamate, Sodium Hydroxide, Phytosphingosine, Ceramide NP, Xanthan Gum, Centella Asiatica Leaf Extract, Ethylhexylglycerin, Bakuchiol, Dehydroacetic Acid, Chlorophyllin-Copper Complex, Rutin, Glycerin, Hydroxycinnamic Acid, Maltodextrin, Sodium Benzoate, Potassium Sorbate.

## Questions or Comments?

Contact [hello@byoma.com](mailto:hello@byoma.com)

## Product Labeling





**BYOMA ACNE TREATMENT SPOT**

salicylic acid paste

| Product Information                                                  |                |                    |               |
|----------------------------------------------------------------------|----------------|--------------------|---------------|
| Product Type                                                         | HUMAN OTC DRUG | Item Code (Source) | NDC:84821-008 |
| Route of Administration                                              | TOPICAL        |                    |               |
|                                                                      |                |                    |               |
| Active Ingredient/Active Moiety                                      |                |                    |               |
| Ingredient Name                                                      |                | Basis of Strength  | Strength      |
| SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ) |                | SALICYLIC ACID     | 20 mg in 1 mL |
|                                                                      |                |                    |               |
| Inactive Ingredients                                                 |                |                    |               |
| Ingredient Name                                                      |                |                    | Strength      |
| CENTELLA ASIATICA LEAF (UNII: 6810070TYD)                            |                |                    |               |
| SODIUM BENZOATE (UNII: OJ245FE5EU)                                   |                |                    |               |
| DEHYDROACETIC ACID (UNII: 2KAG279R6R)                                |                |                    |               |
| SODIUM COPPER CHLOROPHYLLIN (UNII: 1D276TYV9O)                       |                |                    |               |
| HYDROXYCINNAMIC ACID (UNII: IBS9D1EU3J)                              |                |                    |               |
| KAOLIN (UNII: 24H4NWX5CO)                                            |                |                    |               |
| ETHYL LINOLEATE (UNII: MJ2YTT4J8M)                                   |                |                    |               |
| DIATOMACEOUS EARTH (UNII: 2RF6EJ0M85)                                |                |                    |               |
| GLYCERYL STEARATE (UNII: 230OU9XXE4)                                 |                |                    |               |
| BAKUCHIOL (UNII: OT12HJU3AR)                                         |                |                    |               |
| CERAMIDE NP (UNII: 4370DF050B)                                       |                |                    |               |
| ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)                                |                |                    |               |

|                                                     |  |
|-----------------------------------------------------|--|
| <b>WATER</b> (UNII: 059QF0KO0R)                     |  |
| <b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)            |  |
| <b>KOJIC ACID</b> (UNII: 6K23F1TT52)                |  |
| <b>PEG-100 STEARATE</b> (UNII: YD01N1999R)          |  |
| <b>PROPANEDIOL</b> (UNII: 5965N8W85T)               |  |
| <b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)          |  |
| <b>CAMPHOR (NATURAL)</b> (UNII: N20HL7Q941)         |  |
| <b>RUTIN</b> (UNII: 5G06TVY3R7)                     |  |
| <b>BENZYL ALCOHOL</b> (UNII: LKG8494WBH)            |  |
| <b>SODIUM STEAROYL GLUTAMATE</b> (UNII: 65A9F4P024) |  |
| <b>GLYCERIN</b> (UNII: PDC6A3C0OX)                  |  |
| <b>PHYTOSPHINGOSINE</b> (UNII: GIN46U9Q2Q)          |  |
| <b>CETEARYL ALCOHOL</b> (UNII: 2DMT128M1S)          |  |
| <b>SOYBEAN OIL</b> (UNII: 241ATL177A)               |  |
| <b>MALTODEXTRIN</b> (UNII: 7CVR7L4A2D)              |  |
| <b>SAFFLOWER OIL</b> (UNII: 65UEH262IS)             |  |
| <b>POTASSIUM SORBATE</b> (UNII: 1VPU26JZZ4)         |  |
| <b>XANTHAN GUM</b> (UNII: TTV12P4NEE)               |  |

### Packaging

| # | Item Code        | Package Description                                | Marketing Start Date | Marketing End Date |
|---|------------------|----------------------------------------------------|----------------------|--------------------|
| 1 | NDC:84821-008-01 | 15 mL in 1 TUBE; Type 0: Not a Combination Product | 05/12/2025           |                    |

### Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|------------------------------------------|----------------------|--------------------|
| OTC Monograph Drug | M006                                     | 05/12/2025           |                    |

**Labeler** - Byoma Ltd (228554984)